IN THE CLAIMS

Please amend the claims as follows:

Claim 1 (Currently Amended): A process for preparing enantiomerically enriched L- α -amino acids or their salts, comprising [[by]] reacting the corresponding 2-ketocarboxylic acid with an ammonium ion donor in the presence of a whole-cell catalyst which comprises comprising a cloned gene encoding a cofactor-dependent amino acid dehydrogenase and a cloned gene encoding an enzyme which that regenerates the cofactor, at a total input of substrate per reaction volume of ≥ 500 mM, with the addition of the substrate being metered such that the stationary concentration of 2-ketocarboxylic acid is less than 500 mM and the external addition of cofactor, based on the total input of substrate, corresponds to < 0.0001 equivalents.

Claim 2 (Currently Amended): The process as claimed in claim 1,

characterized in that wherein

no cofactor is added to the reaction mixture.

Claim 3 (Currently Amended): The process as claimed in claim 1 and/or 2, characterized in that

use is made of wherein the 2 ketocarboxylic acids which is one that will yield an amino acids acid of the general formula (I)

$$H_2N$$
 COOH

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in which R is alkyl, in particular a space-filling branched alkyl group which that exhibits a tertiary C atom and possesses 5-10 C atoms, for example tert-butyl, [[and]] or substituted alkyls alkyl.

Claim 4 (Currently Amended): The process as claimed in one or more of the preceding claims,

characterized in that claim 1, wherein

the substrate is metered [[in]] in accordance with a fed batch process.

Claim 5 (Currently Amended): The process as claimed in one or more of the preceding claims,

characterized in that claim 1, wherein

the 2-ketocarboxylic acid is kept at a maximum stationary concentration of less than 450 mM, very preferably of less than 400 mM.

Claim 6 (Currently Amended): The process as claimed in one or more of the preceding claims,

characterized in that claim 1, wherein

before it is used, the whole-cell catalyst is pretreated such that the permeability of the cell membrane for the substrate and products is increased as compared with the intact system.